

Division of Heart and Vascular Diseases National Heart, Lung, and Blood Institute, NIH U.S. Public Health Service, DHHS

SPECIAL INSTRUCTIONS FOR PREPARATION OF GRANT APPLICATIONS IN RESPONSE TO RFA: HL-03-009

Specialized Centers of Clinically Oriented Research (SCCOR)

in

Cardiac Dysfunction and Disease

November 2002

This document is also available as a PDF file [92K] (NOTE: only links to external documents will work)

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INSTRUCTIONS FOR THE PREPARATION OF A SCCOR GRANT APPLICATION

A. Introduction

These instructions provide information needed for preparation of an application for a Specialized Center of Clinically Oriented Research (SCCOR) in Cardiac Dysfunction and Disease. They are intended as a companion to RFA: HL-03-009. This RFA appeared in the NIH Guide to Grants and Contracts on November, 2002. The application receipt date for this RFA is September 11, 2003.

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B. General Information

General instructions for preparation of NIH grant applications are contained in the standard NIH grant application kit (PHS Form 398, revised 5/01 and updated 1/25/02, http://grants2.nih.gov/grants/funding/phs398/phs398.html) which is usually available at an applicant institution's business office. Even though the application kit is intended primarily for the regular research-project grant, some of the *general* instructions and forms in it also apply to SCCOR grant applications. However, SCCOR grants have special characteristics. Accordingly, the *specific* instructions in this document were prepared for use with the grant application kit. If questions arise, it is recommended that applicants contact staff of the Division of Heart and Vascular Diseases. To the extent possible, the Division is prepared to discuss plans for developing an application. Inquiries about preparation of SCCOR applications should be addressed to:

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Division of Heart and Vascular Diseases
National Heart, Lung, and Blood Institute
Two Rockledge Centre, Suite 9044
6701 Rockledge Drive - MSC 7940

Bethesda, Maryland 20892-7940 Telephone: (301) 435-0544 Fax: (301) 480-1336

Email: FakundiJ@nhlbi.nih.gov

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C. Specific Instructions for Specialized Centers of Clinically Oriented Research (SCCOR) Applications

- 1. FACE PAGE (Form Page 1)*
 - Type "NHLBI SCCOR" in the top left-hand corner of the face page above the words "GRANT APPLICATION." Enter in Item 1 the title SCCOR in Cardiac Dysfunction and Disease. In Item 2, check "YES" and insert Number: RFA-HL-03-009 and Title: Specialized Center of Clinically Oriented Research. In item 3, indicate the name, degree(s), and position (or equivalent) title of the SCCOR Director. Complete the rest of the page according to the instructions in the application kit, including the signature of the appropriate institution officials. Be sure to attach the *RFA label*, found in the application kit, to the bottom of the face page on the original application. Failure to use the RFA label could result in delayed processing of your application, such that it may not reach the review committee in sufficient time for review.

*Refers to appropriate form in the NIH grant application kit (PHS 398, revised 05/01).

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- 2. DESCRIPTION, PERFORMANCE SITES, AND KEY PERSONNEL (Form Page 2)
 - Provide a brief description of the proposed SCCOR in the space provided. Fill in the performance site(s) and the key professional personnel on this form.

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3. RESEARCH GRANT TABLE OF CONTENTS

• Do not use Form Page 3 provided in the application kit. Instead, prepare a table of contents that identifies by page number all major parts of the SCCOR so that they can be readily located. When listing individual projects and core(s), identify each by a number, title, and responsible investigator in the order in which they appear in the application. Pages should be numbered sequentially throughout the entire application.

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4. DETAILED BUDGET FOR INITIAL BUDGET PERIOD (Form Page 4)

- A detailed budget will be required for EACH PROJECT AND EACH CORE UNIT in their respective sections of the application. (See later under Research Plan)
- To aid in the review of your application, it is suggested that you incorporate a detailed budget for all requested support for the first year (Sample Table 1 in the Illustrations). For each category, such as "Personnel," "Equipment," "Travel," and "Other Expenses," give the amount requested for **EACH PROJECT AND CORE UNIT** with subtotals. If contractual arrangements or "purchased services" involving other institutions or organizations are anticipated (as in project 5 in Sample Table 1 in the Illustrations), include total (direct and indirect) costs associated with such third-party participation in the "Consortium/Contractual Costs" category. In addition, a complete budget for a consortium project is to be developed and identified as such. The published policy governing consortia should be available in the business office of the grantee institution. If clarification is required, Mr. Anthony Agresti at the Grants Operations Branch, Division of Extramural Affairs, NHLBI, should be contacted at (301) 435-0171.

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5. BUDGET FOR ENTIRE PROPOSED PERIOD (Form Page 5)

• Use Form Page 5 in the application kit to show the total SCCOR budget by category requested for each of the 5 years. Justifications for increases in succeeding years should *not* be included here; they should be delineated in the *detailed* budget for *individual* projects.

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6. BIOGRAPHICAL SKETCH (Biographical Sketch Format Page)

• Prepare biographical sketches as described in the application kit; they should be placed in alphabetical order immediately after the summary budget information. Biographical sketches are required for all professional personnel participating in the individual SCCOR projects and core(s). A form entitled "Biographical Sketch" is provided in the application kit.

INFORMATION (Other Support Format Page) See the PHS Form 398 Instructions, page 14.

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7. RESOURCES

• See section 8.b. below, Institutional Environment and Resources.

8. RESEARCH PLAN

- Each project should follow the format for the PHS Form 398 and provide supplementary information when necessary for each section as indicated below. Describe each research project and each core unit in the same detail required for an individual research project grant application, so that the scientific merit can be judged on the basis of the written proposal. Keep in mind that the proposal will be reviewed by experts who can judge, collectively, all areas represented in the proposal but who may not be familiar, individually, with each area of research proposed. Therefore, the description of a project should be concise yet explicit enough to enable experts in related areas to understand the main thrust of each project.
- Page limitations specified for individual (R01) grant applications in PHS Form 398 apply to each project and core unit section. Additional information concerning collaboration and integration between projects and cores and the contribution of each component to the program's specific goals should be succinct. It is the responsibility of the principal investigator to ensure that each component is presented as succinctly as possible. Unnecessarily long, wordy, or confusing presentations are usually perceived as indicative of premature or poorly planned research. The bibliography is not counted toward the 25-page limitation per project.
- The following sections should precede the individual projects.
 - a. Program Introduction and Statement of Objectives
 - A SCCOR should be viewed as a group of interrelated research projects, each of which is not only individually meritorious scientifically but is also complementary to the other projects in the research program and contributes to the integrating theme. The theme of the proposed SCCOR should be established in the first few sentences of the general introduction.
 - Describe the rationale for the proposed total research program. Explain the strategy for achieving the objectives of the overall program and how each research project and core unit relates to that strategy.
 - The general introduction of the overall program description is the appropriate place to indicate any prior collaborative arrangements between investigators in the group, to emphasize the events that have led to the current application, to anticipated unique advantages that would be gained by the research within the proposed SCCOR, to describe how the projects are synergistic and mutually reinforcing, and to explain how the projects collectively would enhance the stated objective of the proposed research.

o In a Table, list each of the research projects and indicate which projects are "clinical research" projects as defined in the RFA.

b. Institutional Environment and Resources

Briefly describe the features of the institutional environment that are relevant to the effective implementation of the overall program. As appropriate, describe available resources such as clinical and laboratory facilities, participating and affiliated units, patient populations, geographic distribution of space and personnel, and consultative resources. The information requested here supplants the "Resources" page in the PHS Form 398 grant application, which is NOT to be used in describing the overall SCCOR program.

c. Organizational and Administrative Structure of a SCCOR

- Describe in detail, and by diagram, if appropriate, the chain of responsibility for decision-making and administration, beginning at the level of SCCOR Director and including investigators responsible for the direction of the research projects and core units. The Director of the overall SCCOR is called the principal investigator; the leaders of the individual projects in a SCCOR are project leaders. Indicate where, in the chain of responsibility, advisory groups (internal and external consultants) would be used.
- Advisory groups: Provide a description of the function of advisory groups (internal and external) used to provide advice and quality control. Identify the expertise needed but do not identify by name the external consultants serving on the advisory groups.
- Specific managerial responsibilities: Indicate who would be responsible for assisting the SCCOR Director with the day-to-day administrative details, program coordination, and the planning and evaluation of the program.
- Relation of the SCCOR organization to the administration of the applicant institution: Describe the relation between the proposed SCCOR and other existing research, academic, and administrative units of the applicant institution such as schools, centers, institutions, departments, and central administration.
- Consortium arrangements: If a SCCOR application includes research
 activities that involve institutions other than the sponsoring organization,
 the program is considered a consortium effort. Such activities may be
 included in a SCCOR grant application, but it is imperative that a

consortium application be prepared so that the programmatic, fiscal, and administrative considerations are explained fully. The published policy governing consortia is available in the business offices of institutions that are eligible to receive Federal grants-in-aid. Consult the latest published policy governing consortia before developing the application. If clarification of the policy is needed, contact Mr. Anthony Agresti at the Grants Operations Branch, Division of Extramural Affairs, NHLBI, at 301-435-0171. Principal Investigators of SCCORs should exercise great diligence in preserving the interactions of the participants and the integration of the consortium projects with those of the parent institution.

Designation of replacement for SCCOR DIRECTOR: Describe the
procedure for appointing a replacement for the SCCOR Director, should
the need arise. The proposed replacement SCCOR Director should meet
all of the applicable criteria. The NHLBI must approve the replacement of
the SCCOR Director should the need arise.

Individual Projects

- 1. Title and number each research project so that it can be readily distinguished from any other project in the program. An individual ABSTRACT should be prepared for each project in the SCCOR as would be required for an R01 (individual Research Grant). The title must NOT exceed 56 typewriter characters/spaces. DO NOT provide a face page (i.e., PHS Form 398 face page) for individual projects. Provide the name, and academic title of the project leader and each participating investigator.
- 2. The budget for each research project must be presented according to the instructions indicated for PHS Form 398. A detailed budget is required for the first and all subsequent years. Include budget requests and explicit detailed budget justifications for all years.
- 3. Research Plan: State the overall objective of the proposed research and explain the relation of the research project to the central theme of the SCCOR and how the project relates to, and both complements and supplements, the other research projects and core units in the program. For projects considered to be "clinical research" projects, please explain the rationale for this designation.

a. Specific Aims

 List the specific aims of the research project for the total period of requested support. Indicate the general priority of each aim in the overall research plan.

b. Background and Significance

 Review the most relevant previous work and describe the current status of research in this field; document with complete references. Specify the overall biomedical significance of the work proposed and the relevance of the research project to the theme of the SCCOR.

c. Preliminary Studies

- Refer to PHS Form 398 Instructions for Preliminary Studies.
- See under Progress Report for Competing Continuation Applications.

d. Research Design and Methods

• Give details of the research plan, including a description of the experiment or other work proposed; present the methods and techniques to be used; note the limitations, if any, of the procedures proposed. Insofar as possible, describe the experiments in the sequence in which they would be conducted. (It is important to convey to the reader that the proposed effort would require the time requested for the project period.)

e. Human Subjects

The NIH policy is that applicants for NIH clinical research grants and cooperative agreements are required to include minorities and women in study populations so that research findings can be of benefit to all persons at risk of the disease, disorder, or condition under study. Special emphasis should be placed on the need for inclusion of minorities and women in studies of diseases, disorders, and conditions that disproportionately affect them. This policy is intended to apply to males and females of all ages. If women or minorities are excluded or inadequately represented in clinical research, particularly in proposed population-based studies, a clear and compelling rationale should be provided. The policy

can be found in the <u>NIH Guide for Grants and Contracts of March 18, 1994</u>, Volume 23, Number 11 and available on the web at: http://grants.nih.gov/grants/guide/notice-files/NOT-OD-00-048.html.

- The composition of the proposed study population must be described in terms of gender, racial, and ethnic groups. These issues must be addressed in developing a research design sample size appropriate for the scientific objectives of the study. This information must be included in PHS Form 398 (rev. 5/01) in Section 8 of the Research Plan (parts a through d) AND summarized in Section 8, part e, Human Subjects Research. Applicants are urged to assess carefully the feasibility of including the broadest possible representation of U.S. racial and ethnic minority groups. However, the NIH recognizes that it may not be feasible or appropriate in all research projects to include representation of the full array of US racial and ethnic minority populations (i.e., American Indian or Alaska Native, Asian, Black or African American, Hispanic or Latino, Native Hawaiian or Other Pacific Islander). The Office of Management and Budget Directive No. 15 describes these categories; it can be found at www.whitehouse.gov/OMB/fedreg/ombdir15.html. The rationale for studies of single US racial or ethnic minority population groups should be provided.
- It is the policy of the NIH that children (i.e., individuals under the age of 21) must be included in all human subjects research, conducted or supported by the NIH, unless there are scientific and ethical reasons not to include them. This policy applies to all initial (Type 1) applications submitted for receipt dates after October 1, 1998. All investigators proposing research involving human subjects should read the NIH Policy and Guidelines on the Inclusion of Children as Participants in Research Involving Human Subjects that was published in the NIH Guide for Grants and Contracts, March 6, 1998, and is available at the following URL address: http://grants.nih.gov/grants/guide/notice-files/not98-024.html
- The usual NIH policies concerning research on human subjects also apply. Form 398 should be followed for human subjects information.
- For foreign components, the policy on inclusion of women and children applies fully. Because the definition of minority differs in other countries, the applicant must discuss the relevance of research involving foreign population groups to the US populations, including minorities.

- If the required information is not contained within the application, the application will be returned.
- Include a Data and Safety Monitoring Plan in the application that
 describes the organizational structures and procedures that will be
 employed to ensure the safety of participants and validity and
 integrity of the data according to NIH policy that can be found at
 NIH policy that can be found at
 http://grants.nih.gov/grants/guide/notice-files/not98-084.html
- Certificates of Confidentiality are issued by NHLBI to protect the privacy of research subjects by protecting investigators and institutions from being compelled to release information that could be used to identify subjects with a research project. Certificates of Confidentiality are issued to institutions or universities where the research is conducted. They allow the investigator and others who have access to research records to refuse to disclose identifying information in any civil, criminal, administrative, legislative, or other proceeding, whether at the federal, state, or local level. Identifying information is broadly defined as any item or combination of items in the research data that could lead directly or indirectly to the identification of a research subject. By protecting researchers and institutions from being compelled to disclose information that would identify research participants, Certificates of Confidentiality help achieve the research objectives and promote participation in studies by assuring privacy to subjects.

Please visit the NIH Certificates of Confidentiality Kiosk (online at http://grants1.nih.gov/grants/policy/coc/index.htm) for additional information including application instructions and frequently asked questions. For further information about Certificates of Confidentiality, to submit an application, or to request an extension or amendment for an existing Certificate awarded by NHLBI, contact:

Mr. David L. Whitmer NHLBI Certificate of Confidentiality Coordinator Chief, Management Policy and Administrative Services Branch 31 Center Drive, Room 5A33, MSC 2490 Bethesda, Maryland 20852-2490

TEL: 301-496-5931 FAX: 301-402-0299

E-MAIL: dave_whitmer@nih.gov

f. Vertebrate Animals (see PHS Form 398 Instructions)

If animals are involved, indicate what kinds are to be used and whether nonhuman primates are to be used, listing the special justifications for their use, and indicate all details for the care, use, treatment, and dispatch of all animals.

g. Literature Cited

• Include a complete citation for each reference in the text.

h. Consortium/Contractual Arrangements and/or Collaborative Arrangements (See PHS Form 398 Instructions)

 Describe the collaboration of investigators within the SCCOR. Describe in detail any other collaborative arrangements anticipated, either within the applicant institution or between institutions.

i. Consultants (see PHS Form 398 Instructions)

 Provide a justification for consultants and attach letters from all individuals confirming their roles in the project.

k. Facilities Available

Describe the facilities to be used by EACH research project. This
is to be accomplished as described under "Resources," in PHS
Form 398 grant application, page 15.

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9. APPENDIX

Appendix material should NOT be included as part of the grant application. Once the application has been received, the Scientific Review Administrator will send the applicant instructions for submitting appendix material. This material includes reprints pertinent to each project or core, and color or half-tone figures. The reviewers will not receive an original copy of the grant application so the appendix should include either originals of the pages or full size copies of any figures where color or clarity is critical to the review. As the PHS Form 398 instructions indicate, the appendix should not be used to circumvent the page limitations.

10. FORMAT OF A CORE UNIT

• A core unit is defined as a resource for the program project that provides centralized services to two or more of the research projects.

a. Title of core unit

 Title and assign a LETTER designation to each core so that it can be readily distinguished from any other core unit. Do NOT exceed 56 typewriter characters/spaces for its title.

b. Names and titles of investigators

• Provide the name and academic title of the core unit leader and each participating investigator.

c. Description of core unit

Describe the function of the core unit as a resource to the SCCOR. This section must present clearly the facilities, techniques, and professional skills that the core unit would provide. As justification for the core unit, indicate briefly the specific research projects that would use the resources of the core unit. A core unit is principally designed as a service or resource component; it would be highly unusual to include research in a core unit (a possible exception would be methodology development). Please contact the Institute staff if you require guidance on this issue.

d. Facilities available

 Describe the facilities to be used by EACH core unit. This is to be accomplished by completing the "Resources" page included in the PHS Form 398 grant application, page 15.

e. Budget for core unit

 Present the budget for each core unit in the format and according to the instructions indicated for Form 398. A budget for the entire proposed project period is required for all subsequent years of support (direct costs only). Include explicit and detailed budget justifications for all years.

f. Relation of core units to research projects

o To aid in the review of your application it is recommended that you prepare in tabular form information concerning the research projects that each core unit would serve and the proportion of the cost of the core unit associated with each research project involved, according to the format presented in Sample Table 2 in the Illustrations.

g. Illustrations in grant application

Sample Table 1 is found in all Form 398 applications (Form Page 4).
 Preparation of Sample Table 2 is suggested to help the reviewers in the evaluation of a SCCOR grant application.

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TABLE 1

Detailed Budget for Initial Budget Period Direct Costs Only

Budget Period ,	From:

Through:

Personnel (Applicant organization only)

For Dollar

Amount Requested, omit cents

		Time/Effort % (of 40		Fringe		
Name	Title of Position	hours per week)	Salary (in dollars)	Benefits (in dollars)	Totals (in dollars)	
Project 1		30,000	3,000	33,000		
Project 2			20,000	2,000	22,000	
Project 3			25,000	2,500	27,500	
Project 4			15,000	1,500	16,500	
Core Unit A			22,000	2,200	24,200	
Core Unit B			10,000	1,000	11,000	
Subtotals			122,000	12,200	134,000	
Consultant Costs (See instructions) Project 2 (\$1,000), Core Unit A (\$2,000) 3,000						
Equipment (Itemize) Project 1 (\$4,000), Project 2 (\$2,000), Project 3 (\$3,000), Project 4 (\$5,000), Core Unit A (\$16,500), Core Unit B 32,500						

(\$2,000)	
Supplies (Itemize by project/core) Project 1 (\$1,000), Project 2 (\$2,000), Project 3 (\$11,500), Project 4 (\$8,900), Core Unit A. (\$16,400), Core Unit B (\$18,200)	79,900
Travel: Domestic	3,000
Travel: Foreign	1,000
Patient Care Cost: Inpatient (Specify Project) Project 4	20,000
Patient Care Cost: Outpatient (Specify Project)	
Alterations and Rennovations (Itemize by project/core)	
Other Expenses (Itemize by project/core) Project 1 (\$1,000), Project 2 (\$1,500), Project 3 (\$1,900) Project 4 (\$1,100), Core Unit A (\$1,500), Core Unit B	
(\$2,200)	9,200
Subtotal Direct Costs for Initial Budget Period	279,800
Consortium/Contractual Costs (See instructions) Project 5 - Collaborating Institution (\$45,200 DC, plus \$10,000 F&A Costs)	
	55,200
Total Direct Costs (Also enter on Page 1, item 7)	335,000

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TABLE 2
Relation of Core Units to Research Projects

Projects	Core Unit A		Core Unit B
Project 1	\$ 3,000	\$ 6,900	
Project 2	13,100	6,000	
Project 3	14,500	10,000	
Project 4	10,000	6,000	
Project 5	20,000	4,500	
Total	\$ 60,600	\$ 33,400	

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